## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

The invention claimed is:

- (Currently amended) A method of generating a cell composition containing cardiomyocytes or cardiomyocyte precursor cells from primate pluripotent stem (pPS) cells obtained from a human blastocyst human embryonic stem (hES) cells, comprising:
  - a) initiating differentiation of the <del>pPS</del> <u>hES</u> cells in suspension culture by forming embryoid bodies;
  - b) culturing the initiated cells so that they differentiate into areas that undergo spontaneous contraction;
    - c) harvesting the differentiated cells;
  - d) separating the harvested cells into fractions according to their based on density; and
  - e) collecting combining the cell fractions containing cells that express cardiac troponin I (cTnI), cardiac troponin T (cTnT), or atrial natriuretic factor (ANF) from an endogenous gene;

thereby generating a cell composition containing cardiomyocytes or cardiomyocyte precursor cells.

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2. (Original) The method of claim 1, wherein the embryoid bodies are plated onto

a surface coated with gelatin or Matrigel®.

3. (Currently amended) The method of claim 1, wherein the cells are differentiated

in the presence of a nucleotide analog that affects DNA methylation; such as 5-

aza-deoxy-cytidine.

4. (Currently amended) The method of claim 1, wherein the cells are differentiated

in a growth environment comprising a morphogen such as activin, and two or

more growth factors.

5. (Original) The method of claim 4, wherein the morphogen is an activin, and the

growth factors include an insulin-like growth factor and a member of the TGFβ

family.

6. (Currently amended) The method of claim 1, wherein the cells are differentiated

in a growth environment containing about 20% serum or serum substitute.

7. (Original) The method of claim 1, wherein the harvested cells are separated by

density centrifugation.

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- 8. (Currently amended) The method of claim 1, wherein the separating comprises distributing cells in the population according to based on their density, and collecting cells at combining cell fractions with a density between ~1.05 and ~1.075 g/mL.
- 9. (Currently amended) The method of claim 1, further comprising culturing the cellected cells combined cell fractions for at least 1 week in a medium containing a compound capable of forming a high energy phosphate bond, an acyl group carrier molecule, and a cardiomyocyte calcium channel modulator.
- 10. (Currently amended) The method of claim 9, further comprising culturing the collected cells combined cell fractions for at least 1 week in a medium containing creatine, carnitine, or taurine.

11.-16. (Canceled)